## **CLEAN VERSION OF THE AMENDED CLAIMS**

## 1-18. (Cancelled)

- 19. (Currently amended) A method for prevention of chronic refractory graft rejection in a human lung transplant recipient comprising administering to the recipient, prior to the development of symptoms associated with transplant rejection, (i) an aerosolized composition comprising a therapeutic dose of cyclosporine and (ii) an effective amount of one or more other immunosuppressive agent.
- 20. (Currently amended) The method of claim 19, wherein the cyclosporine is administered as a dry powder in combination with a propellant.
- 21. (Currently amended) The method of claim 19, wherein the cyclosporine is dissolved in an organic solvent.
- 22. (Previously presented) The method of claim 19, 20 or 21 wherein the dose of cyclosporine is sufficient to achieve deposition levels ranging between 15 and 30 mg in a lung.
  - 23. (Cancelled)
- 24. (Currently amended) The method of claim 19, 20 or 21 wherein the aerosolized composition is co-administered with an anti-inflammatory reagent.
- 25. (Currently amended) A method for ameliorating pulmonary inflammation in a subject having a lung disorder selected from the group consisting of asthma, sarcoidosis, emphysema, cystic fibrosis, idiopathic pulmonary fibrosis, chronic bronchitis, hypersensitivity pneumonitis and eosinophilic pneumonia, comprising administering to the subject an aerosolized

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composition comprising a dose of cyclosporine dissolved in an organic solvent, in an amount effective to inhibit or ameliorate pulmonary inflammation.

- 26. (Currently amended) A method for ameliorating pulmonary inflammation in a subject comprising administering to the subject an aerosolized composition comprising:
- (i) a dose of cyclosporine in dry powder form in an amount effective to inhibit or ameliorate pulmonary inflammation; and
  - (ii) a propellant.
  - 27. (Cancelled)
  - 28. (Cancelled)
- 29. (Currently amended) The method of claim 25 or 26 wherein the dose of cyclosporine is sufficient to achieve deposition levels ranging between 5 and 30 mg in a lung.
- 30. (Currently amended) A method for prevention of graft rejection in a non-lung transplant recipient comprising administering to the non-lung transplant recipient, prior to the development of symptoms associated with transplant rejection, an aerosolized composition comprising a dose of cyclosporine dissolved in an organic solvent, in an amount effective to prevent graft rejection.
- 31. (Currently amended) A method for prevention of graft rejection in a non-lung transplant recipient comprising administering to the non-lung transplant recipient, prior to the development of symptoms associated with transplant rejection, an aerosolized composition comprising:
  - (i) a dose of cyclosporine in dry powder form in an amount effective to

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prevent graft rejection; and

- (ii) a propellant.
- 32. (Cancelled)
- 33. (Currently amended) A method for inhibiting the immune response associated with a T-cell mediated immune disorder in a subject, where said disorder is selected from the group consisting of type IV cell-mediated hypersensitivity, systemic lupus erythematosis, myasthenia gravis, Grave's disease, Hashimoto's thyroiditis, rheumatoid arthritis, scleroderma, and pernicious anemia, comprising administering, to the subject, an aerosolized composition comprising a dose of cyclosporine dissolved in an organic solvent, in an amount effective to inhibit the immune response associated with a T-cell mediated immune disorder.
- 34. (Currently amended) The method of claim 30 or 31 wherein the dose of cyclosporine is sufficient to achieve circulating levels ranging between 50-250 ng/ml.
- 35. (Currently amended) The method of claim 30 or 31 wherein the aerosolized composition is co-administered with a second immunosuppressive agent.
- 36. (Currently amended) A method for inhibiting the immune response associated with a T-cell mediated immune disorder in a subject subject, where said disorder is selected from the group consisting of type IV cell-mediated hypersensitivity, systemic lupus erythematosis, myasthenia gravis, Grave's disease, Hashimoto's thyroiditis, rheumatoid arthritis, scleroderma, and pernicious anemia, comprising administering to the subject an aerosolized composition comprising:
- (i) a dose of cyclosporine in dry powder form in an amount effective to inhibit the immune response associated with a T-cell mediated immune disorder; and

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(ii) a propellant.

37-48. (Cancelled)

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